

K032066 SONOSURG SYSTEMOct 3, 2003
92 days to decisionK032066 · Product code: **LFL** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k032066/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jul 3, 2003
Decision date	Oct 3, 2003
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	The Olympus Optical Co.
Location	Melville, NY, US
Contact	TINA STEFFANIE-OAK
510(k) history	22 submissions · 22 cleared · 1998-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032066/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026